DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
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December 19,2014

Zibo Huaqi Trading Company, Limited
C/O Mr. Ray Wang
Official Correspondent
Beijing Believe Tech. Service Company, Limited
1-202, Build 3, Beijing New World, No.5 Chaoyang Rd.,
Chaoyang District, Beijing, 100024
CHINA

Re: K142508
Trade/Device Name: Powder Free Yellow Vinyl Patient Examination Gloves
Regulation Number: 21 CFR 880.6250
Regulation Name: Patient examination glove
Regulatory Class: I
Product Code: LYZ
Dated: November 4, 2014
Received: November 7, 2014

Dear Mr. Ray Wang:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be

found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Division of Industry and Consumer Education at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to http://www.fda.gov/MedicalDevices/Safety/ReportaProblem/default.htm for the CDRH's Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

You may obtain other general information on your responsibilities under the Act from the Division of Industry and Consumer Education at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm.

Sincerely yours,

Susan Runner DDS, MA

Erin I. Keith, M.S.
Director
Division of Anesthesiology, General Hospital,
Respiratory, Infection Control and Dental Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Food and Drug Administration

Indications for Use

Form Approved: OMB No. 0910-0120
Expiration Date: January 31, 2017
See PRA Statement below.

510(k) Number *(if known)*
K142508

Device Name
Powder Free Yellow Vinyl Patient Examination Gloves

Indications for Use *(Describe)*

The Powder Free Yellow Vinyl Patient Examination Gloves is a disposable device intended for medical purposes that is worn on the examiner's hands or finger to prevent contamination between patient and examiner.

Type of Use *(Select one or both, as applicable)*

☐ Prescription Use (Part 21 CFR 801 Subpart D) ☒ Over-The-Counter Use (21 CFR 801 Subpart C)

CONTINUE ON A SEPARATE PAGE IF NEEDED.

This section applies only to requirements of the Paperwork Reduction Act of 1995.

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Exhibit #7 510(k) Summary

This 510(k) Summary of 510(k) safety and effectiveness information is being submitted in accordance with requirements of SMDA 1990 and 21 CFR 807.92.

The assigned 510(k) Number: _____

1. Date of Preparation: Sept. 8, 2014

2. Sponsor

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3. Submission Correspondent

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4. Proposed Device Identification

Trade Name: Powder Free Yellow Vinyl Patient Examination Gloves
Device Name: Vinyl Patient Examination Gloves (Powder Free)
Common Name: Patient Examination Gloves

Classification: I
Product Code: LYZ
Regulation Number: 21 CFR 880.6250
Review Panel: General Hospital

Intended Use Statement:

The Powder Free Yellow Vinyl Patient Examination Gloves is a disposable device intended for medical purposes that is worn on the examiner's hands or finger to prevent contamination between patient and examiner.

5. Predicate Device Identification

510(k) Number: k091431
Product Name: Patient Vinyl Examination Gloves, Yellow, Powder free
Manufacturer: Zibo Yusheng Product Co., Ltd.

6. Device Description

The proposed device, Powder Free Yellow Vinyl Patient Examination Gloves is a disposable device intended for medical purposes that is worn on the examiner's hands or finger to prevent contamination between patient and examiner.

The proposed is Powder Free Yellow Vinyl Patient Examination Gloves, and includes variations of different size.

7. Non-Clinical Test Conclusion

Bench tests were conducted to verify that the proposed device met all design specifications as was Substantially Equivalent (SE) to the predicate device. The test results demonstrated that the proposed device complies with the following standards:

ASTM D5250-06, Standard Specification for Poly(vinyl chloride) Gloves for Medical Application.
ASTM D 5151-06 (Reapproved 2011), Standard Test Method for Detection of Holes in Medical Gloves.
ASTM D6124-06 (Reaffirmation 2011), Standard Test Method for Residual Powder on Medical Gloves.
ISO 2859-1:1999, "Sampling Procedures for Inspection by Attributes – Part I: Sampling Plans Indexed by Acceptable Quality Level (AQL) for Lot-by-Lot Inspection.
ISO 10993-10: 2010, Biological evaluation of medical devices - Part 10: Tests for irritation and skin sensitization.

8. Substantially Equivalent Comparison Conclusion

Table 1 General Comparison

ITEM	Proposed Device Powder Free Yellow Vinyl Patient Examination Gloves	Predicate Device (k091431) Patient Vinyl Examination Gloves	Remark
Product Code	LYZ	LYZ	SE
Regulation No.	21 CFR 880.6250	21 CFR 880.6250	SE
Class	I	I	SE
Intended Use	The Powder Free Yellow Vinyl Patient Examination Gloves is a disposable device intended for medical purposes that is worn on the examiner's hands or finger to prevent contamination between patient and examiner.	A patient examination glove is a disposable device intended for medical purposes that is worn on the examiner's hands or finger to prevent contamination between patient and examiner.	SE
Powdered or Powered free	Powdered free	Powdered free	SE

Table 2 Device Dimensions Comparison

Proposed Device Powder Free Yellow Vinyl Patient Examination Gloves	Designation	Size				Tolerance
		S	M	L	XL	
	Length, mm	240	240	240	240	min
	Width, mm	85	95	105	115	±5
	Thickness, mm:					
	Finger	0.10				min
	Palm	0.08				min
	Cuff	0.06				min
Predicate Device (k091431) Patient Vinyl Examination Gloves	Designation	Size				Tolerance
		S	M	L	XL	
	Length, mm	240	240	240	240	min
	Width, mm	85	95	105	115	±5
	Thickness, mm:					
	Finger	0.10				min
	Palm	0.08				min
	Cuff	0.06				min
Remark	SE					

Table 3 Performance Comparison

ITEM			Proposed Device Powder Free Yellow Vinyl Patient Examination Gloves	Predicate Device (k091431) Patient Vinyl Examination Gloves	Remark
Colorant			Yellow	Yellow	SE
Physical Properties	Before Aging	Tensile Strength	13 MPa, min	13 MPa, min	SE
		Ultimate Elongation	400 % min	400 % min	SE
	After Aging	Tensile Strength	13 MPa, min	13 MPa, min	SE
		Ultimate Elongation	400 % min	400 % min	SE
	Comply with ASTM D5250			Comply with ASTM D5250	SE
Freedom from Holes			Be free from holes when tested in accordance with ASTM D5151	Be free from holes when tested in accordance with ASTM D5151	SE
Powder Content			0.61 mg per glove	Meet the requirements of ASTM 5250	SE

Table 4 Safety Comparison

ITEM		Proposed Device Powder Free Yellow Vinyl Patient Examination Gloves	Predicate Device (k091431) Patient Vinyl Examination Gloves	Remark
Material		Vinyl	Vinyl	SE
Biocompatibility	Irritation	Under the conditions of the study, not an irritant	Comply with ISO 10993-10	SE
	Sensitization	Under conditions of the study, not a sensitizer		
Label and Labeling		Meet FDA's Requirements	Meet FDA's Requirements	SE

The proposed device, Powder Free Yellow Vinyl Patient Examination Gloves, is determined to be Substantially Equivalent (SE) to the predicate device in respect of safety and effectiveness.